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The Influence of Hydrogen on Thermal and Catalytic Cracking of n-Octane

69791 \$/055/59/000/06/20/027 B004/B002

(Table 1, Fig.2). The reaction furnace No. 1 used first had too much of lost space (gaps not filled by the catalyst) in which thermal cracking took place due to overheating. By using reaction furnace No. 2 thermal cracking of octane could be reduced to about one half. Table 2 and Fig. 3 give the results of the reaction after the addition of hydrogen and nitrogen. Hydrogen increases the yield of thermal cracking by 6%, and nitrogen by 3%. Fig. 4 shows that the yield of thermal cracking at 500° increases up to a constant value if the molecular ratio of H<sub>2</sub>: C<sub>8</sub>H<sub>18</sub> is increased. Fig. 5 shows the same result at 530°. The yield of catalytic cracking was not affected by hydrogen. Table 3 gives the analyses of the cracking products. In the presence of hydrogen, isomerization of n-octane set in. At 500° 5% of 3-methylheptane was obtained and at 550° 10%. The authors mentioned B. T. Abayeva (Ref. 4). There are 5 figures, 3 tables, and 11 references, 6 of which are Soviet.

ASSOCIATION: Kafedra fizicheskoy khimii (Chair of Physical Chemistry)

SUBMITTED: February 25, 1959

Card 2/2

TOPCHIYEVA, K.V.; MOSKOVSKAYA, I.F.; BODROVA, L.G.; KRUPENYA, E.I.

Studying the nature of the activity of aluminosilicate catalysts. Vest Mosk. un. Ser. mat., mekh., astron., fiz., khim. 14 no.2: 225-235 '59 (MIRA 13:3)

 Kafedra fizicheskcy khimii Moskovskogo gosuniversiteta. (Catalysts) (Aluminosilicates)

5(4)

AUTHORS: Smirnova, I. V., Topchiyeva, K. V., Smetanko, N. P. (Moscow)

TITLE: The Adsorption From Solutions of Alkylaromatic Hydrocarbons on Industrial Catalysts 2. (Adsorbtsiya iz rastvorov alkil-

aromaticheskikh uglevodorodovnapromyshlennykh katalizatorakh.2)

PERIODICAL: Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 5. pp 1059 - 1064 (USSR)

ABSTRACT: This paper shows the results of the investigation of the adsorption of allyl benzene, propenyl benzene, and - in comparison - n-propyl benzene from solutions of n-heptane on Al<sub>2</sub>O<sub>3</sub> at 20° and 40°. Table 1 shows the physical data

of the hydrocarbons used. Figure 1 shows the isothermal adsorption lines at 20°, figure 2 at 40°. The absolute isothermal adsorption lines and their molecular constants were determined considering the extent of the specific surface of Al<sub>2</sub>O<sub>3</sub>. Figure 2 shows the isothermal lines, table 2 the data

obtained. The thickness of the adsorption layer of propenyl benzene agrees with the theoretically calculated thickness

Card 1/2 of the benzene ring = 3.7 Å. Thus the molecules of propenyl

The Adsorption From Solutions of Alkylaromatic Hydrocarbons on Industrial Catalysts 2.

sov/76-33-5-16/33

benzene show a parallel orientation towards the catalyst surface with the surface of the benzene ring. The same is true of allyl benzene and n-propyl benzene. The presence of a double bond in the side chain does not change the planoparallel orientation of the benzene derivative. The adsorbability of the hydrocarbons with various molecular volume decreases in the order propenyl-, allyl-, n-propyl benzene. A conjugated double bond increases the adsorption potential. Adsorption decreases with increasing temperature, the adsorption layers become less dense. There are 3 figures, 2 tables, and 16 references, 14 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova

(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: October 12, 1957

Card 2/2

sov/76-33-7-39/40

5(4) AUTHORS : Gerasimov, Ya. I., Topchiyeva, K. V., Semiokhin, I. A.,

TITLE:

Georgiy Mitrofanovich Panchenkov. On the Occasion of His 50th

Birthday

PERIODICAL:

Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 7,

pp 1674 - 1675 (USSR)

ABSTRACT:

On April 24, 1959 G. M. Panchenkov, a well-known Soviet specialist in physical chemistry and Professor at the Moskovskiy institut neftekhimicheskoy i gazovoy promyshlennosti im. I. M. Gubkina and Moskovskiy gosudarstvennyy universitet (Moscow Institute for Petroleum-Chemical and Gas Industry imeni I. M. Gubkin and Moscow State University), celebrated his 50th birthday. The main fields with which he was concerned are the kinetics of heterogeneous catalytic processes, the methods of separating and analyzing isotopes, and the theory of the liquid phase. His investigations of the mechanism of the transformation of hydrocarbons on aluminum silicate catalysts by the use of deuterium as a marking atom as well as his publications on the theory of viscosity are especially worth mentioning. For the latter he was awarded the Stalin Prize for the field of sciences in 1952. The

Card 1/2

CIA-RDP86-00513R001756310011-6" APPROVED FOR RELEASE: 08/31/2001

Georgiy Mitrofanovich Panchenkov. On the Occasion of His 50th Birthday

SOV/76-33-7-39/40

method of separating boron isotopes devised by G. M. Panchenkov et al was demonstrated at the Vsesoyuznaya promyshlennaya vystavka (All-Union Industrial Exposition) and was awarded a diploma of the second class, this method also has won general appreciation at international expositions in Geneva, Leipzig, Peking, and Warsaw. Professor G. M. Panchenkow, who is also a teacher, founded the Kafedra fizicheskoy i kolloidnoy khimii (Chair of Physical and Colloid Chemistry) at the above-mentioned Institute as well as the Laboratoriya khimii i razdeleniya izotopov v MGU (Laboratory for Chemistry and Isotope Separation at Moscow State University), which have been headed by him up to this day. dissertations for the degree of Doctor and 15 dissertations for the degree of Candidate were completed under his supervision. He published 2 monographs, about 100 scientific articles, and obtained 10 patents for his inventions. G. M. Panchenkov is a member of the International Committee for Constants. Furthermore, he was awarded the orders "Krasnaya Zvezda" and "Znak Pocheta" as well as the title of Laureate of the Stalin Prize. There is 1 figure.

Card 2/2

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

sov/20-124-1-38/69 Topchiyeva, K. V., Romanovskiy, B. V. 5(4) AUTHORS: Determination of the Adsorption Coefficients of Ether, Water, and Ethylene by the Kinetic Method (Opredeleniye adsorbtsionnykh koeffitsiyentov efira, vody i etilena kineticheskim metodom) TITLE: Doklady Akademii nauk SSSR, 1959, Vol 124, Nr 1, pp 135-138 PERIODICAL: (USSR) There are two essentially different methods of determining the adsorption coefficients, viz. the kinetic- and the ABSTRACT: adsorption method. However, the authors believe that only the kinetic method shows the proper way of determining the adsorption coefficients. The present paper discusses the method developed by A. V. Frost (Ref 4) for determining adsorption coefficients. For the reactions occurring according to the scheme  $A_1 \rightarrow Y_2 A_2 + Y_3 A_3 + \cdots + Y_i A_i$  he suggested the equation  $v_0 \ln(1/(1-y)) = \alpha + \beta v_0 y$ ;  $v_0$  denotes the rate at which the initial material is introduced into the reactor (expressed in Mol per unit of time and weight of the catalyst); y - the degree of transformation; the constants  $\alpha$  and  $\beta$  do not depend on  $v_0$  and y. A formula for  $\alpha$  is written down. The Card 1/3

Determination of the Adsorption Coefficients of Ether, Water, and Ethylene by the Kinetic Method SOV/20-124-1-38/69

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quantity 1/0 can be written down as linear function of dilution. For the purpose of determining the adsorption coefficient of a substance slowing down catalytic reaction it is necessary to carry out two series of investigation: one with a dilution by the substance the adsorption coefficient of which is to be determined, and another by means of an inert diluent. The present paper deals with the slowing-down influence of additions of ether, water, and ethylene on the dehydration rate ethyl alcohol by way of ammonium oxide in order to determine the adsorption coefficients of these substances. Argon was used as inert diluent. The authors determined the adsorption coefficients of water and ether at 2500 and of ethylene at 430°. At 250° alcohol is decomposed only in water and in ether. With a rise in temperature, the degree of transformation is reduced, and this diminishes the accuracy of the kinetic equation applied. At 430° alcohol is dehydrated to ethylene and water and the ether content in the catalyzed product is extremely low. All investigations were carried out under atmospheric pressure in an ordinary laboratory apparatus. The authors carried out two series of tests: one with dilution

Card 2/3

Determination of the Adsorption Coefficients of Ether, Water, and Ethylene by the Kinetic Method

SOV/20-124-1-38/69

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by water, ether, and argon at  $250^{\circ}$ , and the other with dilution by water, ethylene, and argon at  $430^{\circ}$ . The kinetics is conserved in the investigated interval of dilutions. On the basis of the experimentally obtained values of  $\alpha$  the diagrams for the dependence of  $1/\alpha$  on dilution were constructed. From the slope of the curves the numerical values of the adsorption coefficients were calculated. There are 2 figures, 1 table, and 11 references, 9 of which are Soviet.

ASSOCIATION:

Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova (Moscow State University imeni M. V. Lomonosov)

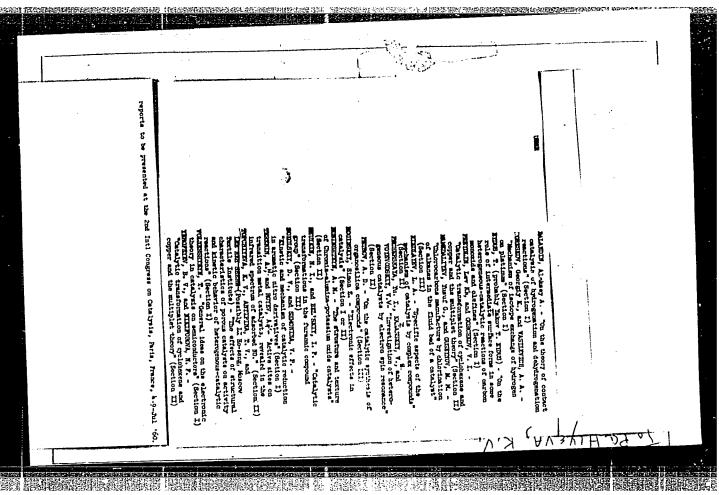
PRESENTED:

August 28, 1958, by A. N. Frumkin, Academician

SUBMITTED:

August 27, 1958

Card 3/3



APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

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TOPCHIYEVA, K.V.; ZEN'KOVICH, I.A.; TRESHCOVA, Ye.G.

Effect of hydrogen on the thermal and catalytic cracking of n-octane. Vest.Mosk.un.Ser.mat.,mekh.,astron.,fiz.,khim.
(MIRA 13:10)
no.6:164-170 159.

1. Kafedra fizicheskoy khimii Moskovskogo universiteta. (Cracking process) (Octane)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

s/195/60/001/003/013/013 B002/B058

AUTHORS:

Topchiyeva, K. V., Antipina, T. V., Li Khe-suyan;

TITLE:

The Effect of the Structural Porosity of Catalysts on Their

Activity and the Kinetic Parameters of the Course of the

Cracking Reaction

PERIODICAL: Kinetika i kataliz, 1960, Vol. 1, No. 3, pp. 471 - 477

TEXT: The effect of the size of pores of an alumino silicate catalyst on the cracking reaction of cumene between 350 and 500°C was studied. A catalyst of the following composition was used: 12% Al203 and 88% SiO2.

The various sizes of pores between 12 and 115 A were obtained by replacing the intermicellar water to a different degree by isobutyl alcohol, isoamyl alcohol or cumene. Moreover, an industrially produced catalyst and a catalyst of the type Gudri (Goodry?) were studied. The structure of the samples was calculated from the adsorption isotherm of methanol vapor at 20°C. The kinetic of the cracking reaction of cumene between 350 and 475°C

Card 1/2

CIA-RDP86-00513R001756310011-6" APPROVED FOR RELEASE: 08/31/2001

The Effect of the Structural Porosity of \$\frac{5}{195}/60/001/003/013/013\$

Catalysts on Their Activity and the Kinetic B002/B058

Parameters of the Course of the Cracking

Reaction

is described very well by the following equation by A. V. Frost:  $v \ln(1/1-y) = \alpha + \beta v_0 y, \quad v \text{ is the volume rate of the addition of the initial material in mmoles/geh; y is the degree of reaction; <math>\alpha$  is the apparent reaction rate constant in mmoles/geh;  $\beta$  is a constant equal to apparent reaction rate constant in mmoles/geh;  $\beta$  is a constant equal to apparent reaction rate constant in mmoles/geh;  $\beta$  is a constant equal to apparent reaction rate constant in mmoles/geh;  $\beta$  is a constant equal to apparent reaction rate considerable by a break of the curve. This range to the diffusion range is manifested by a break of the curve. This corresponds to a considerable change of the activation energy. There are 9 figures, 2 tables and 15 references: 13 Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State

University)

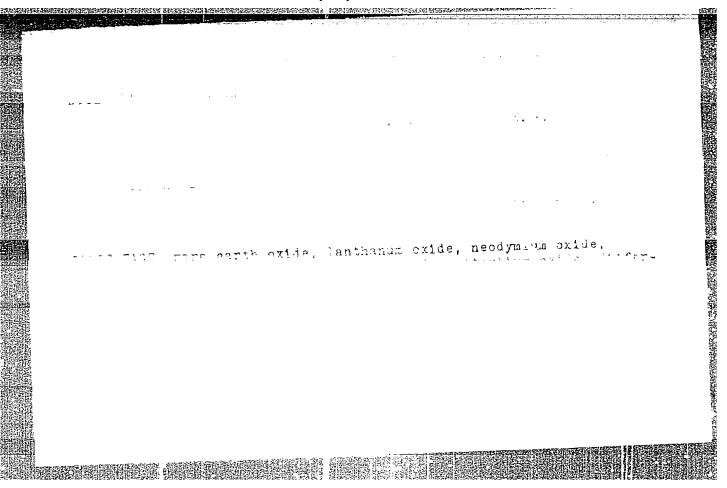
SUBMITTED: March 23, 1960

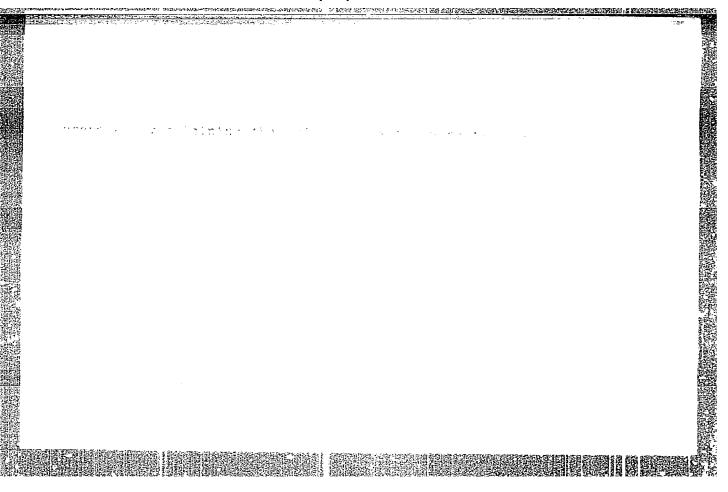
Card 2/2

YEGOROV, M.M.; IGNAT'YEVA, L.A.; KISELEV, V.F.; KRASIL'NIKOV, K.G.; TOPCHIYEVA, K.V.

Surface properties of catalytically active aluminum oxide. Zhur. fiz. khim. 36 no.9:1882-1889 S '62. (MIRA 17:6)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, fizicheskiy fakul'tet i khimicheskiy fakul'tet.



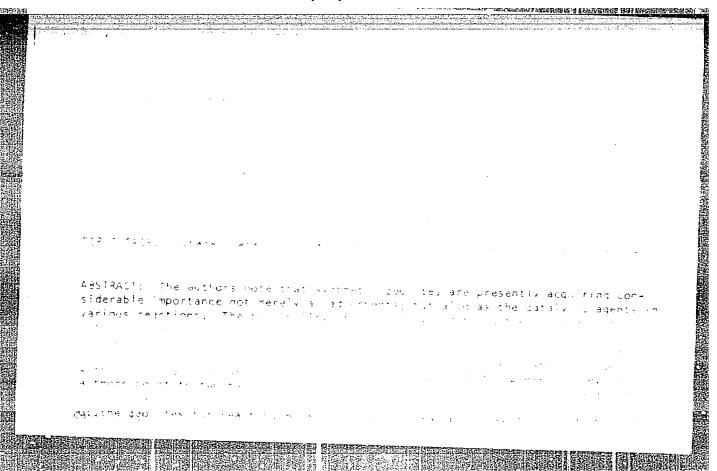


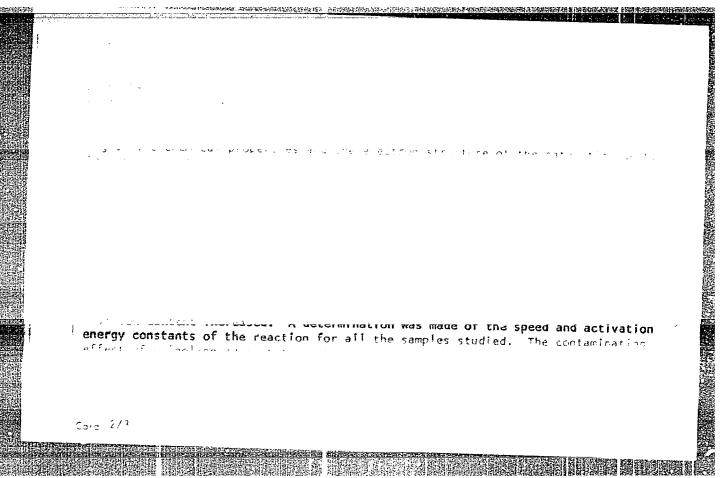
KUBASOV, A.A.; SMIRNOVA, I.V.; TOPCHIYEVA, K.V.

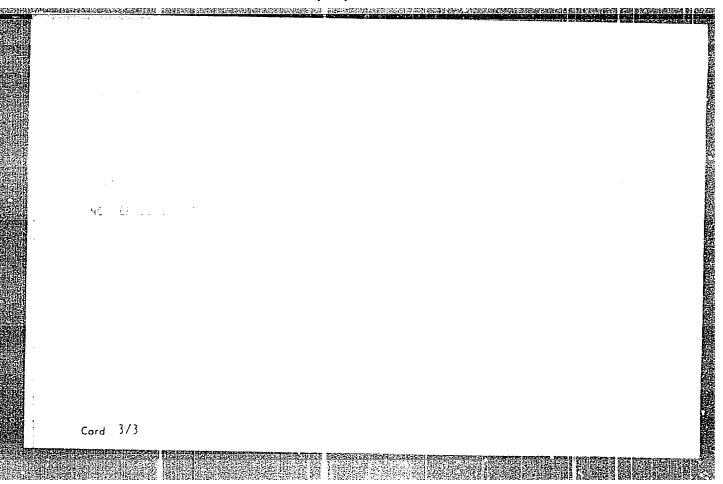
Gas chromatographic determination of the heats of adsorption of hydrocarbons on aluminum exide, Kin. i kat. 5 no.3:520-525 My-Je 164. (MIRA 17:11)

1. Moskovskiy josudarstvennyy universitet imeni Lemonosova, khimicheskiy fakulitet.

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"







SIGNACHA, A.A.; MESEOVSKATA, I.F.; TORGRITIVA, K.V.

Determination of the heats of high temperature adsorption of hydrogarbons on cracking entalysis. Vest. Mask. un. For. 22 Khim. 20 no. 5:13-18 S-0 '65. (MEA 18:12)

1. Kafedra fizichenkoy khimit Moskowskogo gosnahestvennogo universiteta. Submitted June 25, 1965.

ZEN'KOVICH, I.A.; TRESHCHOVA, Ya.G.; TOPCHIYEVA, K.V.

Transformation of phenylcyclopropane on Junioum exide with boron fluoride. Vest. Mosk. un. Ser. 2:Khim. 20 no. 5:19-22 S-0 '65. (NIRA 18:12)

1. Kafedra fizicheskoy khimii Moskovskogo gosudarstvennogo universiteta. Submitted Dec. 15, 1964.

KRASIL'NIKOVA, M.K.; TOPCHIYEVA, K.V.

Chemisorption of ethylene on yttrium oxide. Kin. i kat. 6 no. 6: 1118-1121 N-D 65 (MIRA 19:1)

1. Moskovskiy gosudarstvennyy universitet imeni Iomonosova, khimicheskiy fakul tet. Submitted June 28, 1965.

BORESKOVA, Ye.G.; TOPCHLYEVA, K.V.; PIGUZOVA, L.I.

Catalytic activity of synthetic zeolites in the cracking of cumene. Kin. i kat. 5 no.5:903-909 S-0 '64. (MIRA 17:12)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimicheskiy fakul tet.

TOICHIY EVA, K.V.; MOSKOVSKAYA, I.F.; DOBROKHOTOVA, N.A.

Use of thermometric titration for measuring the acidity of solid oxide catlaysts. Kin. 1 kat. 5 no.5:910-915 S-0 '64.

(MIRA 17:12)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.

L 1681-66 EWT(m)/EPF(c)/EWP(j)/EWP(t)/EWP(b) JD/JG/RM

ACCESSION NR: AP5020989

UR/0195/65/008/004/0751/0751 546-643-31-44

AUTHOR: Topchiyeva, K. V.; Stetsenko, V. Ya. 55

TITLE: New catalytic properties of yttrium oxide

SOURCE: Kinetika i kataliz, v. 6, no. 4, 1965, 751

TOPIC TAGS: catalysis, yttrium compound, hydrogenation, isomerization, ethylene, propylene, acetylene

ABSTRACT: The article presents the results of a series of experiments which point to the sufficiently high catalytic activity of yttrium oxide in hydrogenation and isomerization reactions. Hydrocarbons investigated included ethylene. propylene, divinyl, acetylene, and 2-methyl-pentene. Temperatures varied from 110-320C, the hydrogen/hydrocarbon ratio from 3:1 to 12:1, the space velocity from 144 to 576 hr<sup>-1</sup>, and the total degree of conversion from 70 to 100%. A necessary condition for the appearance of the catalytic activity of yttrium oxide is its activation with hydrogen. A sufficiently constant activity of the yttrium

Card 1/2

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ACCESSION NR: AP5020989												
- Airester amount of hydror	ne and activation rate with hydrogen, with a comp carbon in the reaction mixture. The work indicated ect to poisoning by traces of moisture and oxygen											
ASSOCIATION: Moskovskii gosudarstvennyi universitet im. M. V. Lomonoso Khimicheskii fakulitet! (Moscow State University, Departm												
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GANICHEMEO, L.G.: TOPOR, N.D.; TCHCHIXEVA, K.V.

l. Kafedra fizioneskoy knimii Moskovskogo universiteta.

TOPCHIYEVA, K.V.; ROMANOVSKIY, B.V.

Girculation method used in studying the kinetics of heterogeneous

catalytic reactions. Part 1: Dehydration of ethyl alcohol on aluminum oxide. Kin. i kat. 6 no.2:279-284 Mr-Ap '65. (MIRA 18:7)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimiche-skiy fakul'tet.

TORUNIVEVA, K.V., MCSKOVEKAYA, I.P.: STETSENKO, V.Ya.

Electric conductivity of aluminosilucate cutalyons for precising.

Kin., kat. 5 nc.611026-1033 N.O 46A. (MIRA 18:3)

J. Moskovskiy gosudarstvennyy universitet limeni Loronosova zhimicheskiy fukulitet.

VENTYAMINOV, S.A.; TOPCHIYEVA, K.V.

Gas chromatographic method of studying the adsorption of acetylene and vinyl chloride on technical aluminum oxide. Kin.i kat. 5 no.6:1107-1110 N-D \*\*164. (MIRA 18:3)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimicheskiy fakulitet.

econtrol produced and produced

BONESKOVA, Ye.G., LYGIN, V.L., TORTHIYEVA, K.V.

Infrared spectroscopy study of the nature of active centers in the cracking of cumene catalyzed by decationized realities. Kin. I kat. 5 no.631115-1118 N-D 464. (MIRA 18:3)

1. Moskovskiy gosudarstvennyy universitet imeni lomonosovs, khimicheskiy fakul tet.

SMIRNOVA, I.V.; KUBASOV, A.A.; BYULOV, Martin; TOPCHIYEVA, K.V.

Heats of wetting of aluminum oxide by solutions of methylcyclohexenes in n-heptane. Dokl. AN SSSR 160 no.1:170-173 Ja 165.

(MIRA 18:2)

1. Moskovskiy gosudarstvennyy universitet. Submitted July 2, 1964.

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

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TOPCHIYEVA, K.V.; ROMANOVSKIY, B.V.; BIRYUKCVICH, M.M.

Study of the inhibiting action of cumene hydroperoxide in the cracking reaction. Vest. Mosk. un. Ser. 2: Khim. 18 no.3: 18-23 My-Je \*63. (MIRA 16:6)

l. Kafedra fizicheskoy khimii Moskovskogo universiteta.

TOPCHIYEVA, K. V.; SHARAYEV, O. K.; PEREL'MAN, A. I.; RYABOVA, A. A.

Effect of the porous structure of the aluminosilicate carrier on the polymerizing activity of the chromium oxide catalyst. Plast. massy no. 5:11-13 '64. (MIRA 17:5)

TOPCHIYEVA, K. V.; SMIRNOVA, I. V.; KUBASOV, A. A.

"Concerning the mechanism of cyclene isomerization over alumina."

report submitted to 3rd Intl Cong on Catalysis, Amsterdam, 20-25 Jul 64.

Moscow State Univ im Lomonosov.

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

TOPCHIYEVA, K.V.; MOSKOVSKAYA, I.F.; STETSENKO, V.Ya.

Electric conductivity of synthetic zeolites. Zhur.fiz.khim. 37
no.8:1883-1885 Ag '63. (MIRA 16:9)
(Zeolites--Electric properties)

TOFTSHIYEVA, K. V.

A. P. Ballod and K. V. Toptshiveva

"The Nature of the Catalytic Effect of Aluminumsilicates." Progress of Chemistry 20, 161-175, April 1951, Moscow

ABSTRACT AVAILABLE

D-50054

TOPCHIYEVA, K.V.; VEN'YAMINOV, S.A.

Kinetics of hydrochlorination of acetylene on aluminum oxide. Kin. 1 kat. 4 no.3:450-460 My-Je 163. (MIRA 16:7)

l. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimicheskiy fakul'tet.

(Acetylene) (Hydrochloric acid)

(Aluminum oxide)

L 12593-63 EPF(c)/EWT(E)/BDS Pr-4 RM/WW S/0189/63/000/003/0018/0023

AUTHOR: Topchiyeva, K. V.; Romanovskiy, B. V.; Biryukovich, M. M.

58

TITIE: Astudy of the inhibitory effect of cumenehydroperoxide in the cracking reaction

SOURCE: Moscow. Universitet. Vestnik. Seriya 2. Khimiya, no. 3, 1963, 18-23

TOPIC TAGS: cumene, cumenehydroperoxide, cracking, catalysis of cracking, alumosilicate catalyst, acetophenone, inhibition of catalysis

ABSTRACT: While the cracking of cumene is today widely used in evaluating the capacity of alumosilicate catalysts, it is essential to take into consideration the inhibiting effect of hydroperoxides of cumene on the kinetic laws of the reaction. This seems to be due to competition of cumene and hydroperoxide for the active centers of the catalyst. In this present work the adsorption coefficients of the hydroperoxide of cumene and its decomposition products (acetophenone) and acetone) were determined. The principle of the method used consisted in studying the effect on the reaction rate constant of various amounts of hydroperoxide, as well as using dilutions by an inert substance - cyclohexane. The obtained results showed that the magnitude of the adsorption coefficient of hydroperoxide is very

Card 1/2

	L 12593-63 ACCESSION NR: AP3001602		2	<b>-</b> ,					
	sizable - an indication that the adsorption equilibrium is substantially shifted towards the formation of a stable complex between the hydroperoxide and the catalyst's active centers. It was also found that acetophenone equals the hydroperoxide in inhibition effect, while acetone has a four times lesser adsorption coefficient. Orig. art. has: 4 formulas, 4 charts, and 1 table.  ASSOCIATION: Moskovskiy universitet, kafedra fizicheskoy khimii (Moscow University, Department of Physical Chemistry)								
	SUBMITTED: 11Jul61	DATE ACQ: 09Jul63	ENCL: 00						
	SUB CODE: 00	NO REF SOV: 011	OTHER: 002						
A.S., 13									
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TOPCHIYEVA, K.V.; RAKHOVSKAYA, S.M.; KUCHKAYEVA, I.K.; SHAMINA, I.S.; YURKEVICH, A.A.

Modifications of the supporting structure of phosphoric acid catalysts in the ethylene hydration process. Neftekhimiia 3 no.2:271-275 Mr-Ap '63. (MIRA 16:5)

1. Saratovskiy gosudarstvennyy universitet imeni N.G.Chernyshevskogo, Nauchno-issledovatel'skiy institut khimii, Moskovskoy gosudarstvennyy universitet imeni Lomonosova i Leningradskiy tekhnologicheskiy institut imeni Lensoveta.

(Phosphoric acid) (Ethylene) (Hydration)

TOPCHIYEVA, K.V.; ROMANOVSKIY, B.V.; KHO-SHI TKHOANG

Kinetics of cumene cracking on 10% zeolite. Dokl.AN SSSR 149 no.3:644-647 Mr '63. (MIRA 16:4)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.

Predstavleno akademikom M.M.Dubininym.

(Cumene) (Cracking process) (Zeolites)

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SMIRNOVA, I.V.; KARPUKHINA, G.V.; TOPCHIYEVA, K.V.

Adsorption of allylbenzene and allylcyclohexane on a chromia catalyst. Neftekhimia 3 no.1:71-73 Ja-F 163. (MIRA 16:2)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.
(Benzene) (Cyclohexane) (Adsorption)

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SMIRNOVA, I.V.; TOPCHIYEVA, K.V.; KUBASOV, A.A.; SAVCHENKO, L.V.

Adsorption of methylcyclohexene from solutions at elevated temperature. Dokl. AN SSSR 147 no.3:660-662 N '62. (MIRA 15:12)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosqva. Predstavleno akademikom P.A. Rebinderom. (Cyclohexene) (Adsorption)

ORMANETS, V.; TRONOVA, V.A.; TOPCHIYEVA, K.V.

Simplified method for the determination of mono-,di-, and triethylamines in a six-component mixture obtained in the catalytic deamination of aliphatic amines over dehydrating oxide catalysts. Zhur.anal.khim. 17 no.9:1109-1113 D 162. (MIRA 16:2)

1. M.V. Lomonosov Moscow State University. (Amines)

HINTER CONTROL OF THE CONTROL OF THE

BALANDIN, A.A., akademik, red.; KOEOZEV, N.I., prof., red.; LEBEDEV, V.P., dots., żam. red.; MAL'TSEV, A.N., zam. red.; AGRODOLOV, A.Ye., dots., zam. red.; TOPCHIYEVA, K.V., prof., red.; YUR'YEV, Yu.K., prof., red. PANCHENKOV, G.M., prof., red.; SOKOL'SKIY, D.V., akademik, red.; VOL'KENSHTEYN, F.F., prof., red.; LAZAREVA, L.V., tekhn. red.

[Catalysis in the institutions of higher learning; papers of the First Interuniversity Conference on Catalysis]Kataliz v vysshei shkole; trudy. Moskva, Izd-vo Mosk. univ. No.1. Pt.2. 1962. 325 p. (MMRA 15:10)

1. Mezhvuzovskoye soveshchaniye po katalizu. 1st, 1958. 2. Akademiya nauk Kazakhskoy SSR (for Sokol'skiy). 3. Khimicheskiy fakul'tet Moskovskogo gosudarstvennogo universiteta (for Yur'yev). (Catalysis)

SERGEYEV, G.B.; SHARAYEV, O.K.; TOPCHIYEVA, K.V.; PEREL'MAN, A.I.; TOPCHIYEV, A.V.

Electron paramagnetic resonance studies of chromium oxide catalysts for ethylene polymerization. Neftekhimia 2 no.1:18-20 Ja-F '62. (MIRA 15:5)

1. Institut neftekhimicheskogo sinteza AN SSSR i Khimicheskiy fakul'tet Moskovskogo universiteta.

(Catalysts—Spectra) (Ethylene) (Polymerization)

TOPCHIYEVA, K.V.; ROSOLOVSKAYA, Ye.N.

Effect of the dehydration of an aluminosilicate catalyst on its acidity. Neftekhimiia 2 no.3:298-304 My-Je '62.

(MIRA 15:8)

1. Moskovskiy gosudarstvenny universitet imeni Lomonosova, khimicheskiy fakul'tet.

(Aluminosilicates) (Dehydration (Chemistry))

(Hydrogen-ion concentration)

SHARAYEV, O.K.; TOPCHIYEVA, K.V.; PEREL'MAN, A.I.; TOPCHIYEV, A.V.

Nature of the induction period in the polymerization of ethylene on a chromium oxide catalyst. Neftekhimiia 2 no.2:187-188 Mr-Ap '62. (MIRA 15:6)

1. Institut neftekhimicheskogo sinteza AN SSSR i Moskovskiy universitet, kafedra fizicheskoy khimii.

(Ethylene polymers) (Catalysts, Chromium)

TOPCHIYEVA, K.V.; ROSOLOVSKAYA, Ye.N.

Effect of the heat treatment of aluminosilicate catalysts in a vacuum of their structure. Neftekhimiia 2 no.2:175-178 M5-Ap '62.

(MIRA 15:6)

1. Moskovskiy gosudarstvenny universitet imeni M.V.Lomonosova khimicheskiy fakul'tet.

(Aluminosilicates)

#### "APPROVED FOR RELEASE: 08/31/2001

## CIA-RDP86-00513R001756310011-6

S/204/62/002/001/001/007 I032/I232

**AUTHORS:** 

Sergeyev, G. B., Sharayev, O. K., Topchiyeva, K. V., Perel'man, A. I., and Topchiyev,

A. V.

TITLE:

Investigation of chromic oxide catalysts for polymerisation of ethylene by the method

of electron paramagnetic resonance

PERIODICAL: Neftekhimiya, v. 2, no. 1, 1962, 18-20

TEXT: The aim of this study was the verification of the hypothesis, previously expressed by the authors, that the activity of the catalyst is produced under the action of the reacting substance, ethylene. Experiments on polymerisation of ethylene over chromic oxide catalysts were carried out and the EPR spectra of the catalyst withdrawn from the reaction zone at different stages of the process were taken. The catalyst was prepared by impregnating aluminum silicate with an aqueous solution of chromic anhydride and subsequent activation. Two varieties of the catalyst, differing by the method of activation, were used. One was activated in a current of air at 500°, the other one— under vacuum at 350°. The catalyst activated under vacuum displayed an induction period. The EPR spectra of the two varieties of catalyst, taken at identical stages of the polymerisation process, were found to be practically identical with respect both to the line width and the value of

Card 1/2

Investigation of chromic oxide catalysts...

S/204/62/002/001/001/007 I032/I232

the g factor (which was 1.97). The identity of the active centres in the two varieties of the catalyst was thus established. The observed narrow EPR line is attributed to a compound of quinquevalent chromium and the Cr<sup>5+</sup> ions are considered to constitute the active centres. The induction period in the catalyst activated under vacuum is interpreted as the time necessary for the reduction of Cr<sup>6+</sup> by ethylene. There are 2 figures.

ASSOCIATION: Institut neftekhimicheskogo sinteza AN SSSR, Khimicheskii fakultet Moskovskogo

Universiteta (Institute of Petrochemical Synthesis, AS USSR, Chemistry Faculty, Uni-

versity of Moscow)

SUBMITTED: Novem

November 24, 1961

Card 2/2.

Card 1/3

S/076/62/036/009/002/011 B101/B102

AUTHORS: Tegorov, h. M., Ignat'yeva, L. A., Kiselev, V. F., Krasil'ni-kov, h. G., and Topchiyeva, K. V.

PERIODICAL: Zhurnal fizicheskoy khimii, v. 36, no. 9, 1962, 1882 - 1889

TEXT: The specific heat of wetting of commercial  $\text{Al}_2\text{O}_3$  by water, methanol, ethanol, and n-heptane, and the content of structural water  $\text{Al}_2\text{O}_3$  were measured, the phase composition of  $\text{Al}_2\text{O}_3$  was determined by x-ray analysis, and the infrared spectrum of deuterated  $\text{Al}_2\text{O}_3$  was taken. Whereas with n-heptane the heat of wetting is independent of the content of structural water in  $\text{Al}_2\text{O}_3$ , it increases, in the case of water and alcohols, with inversing thermal dehydration of  $\text{Al}_2\text{O}_3$ . Since, however, the specific surface of  $\text{Al}_2\text{O}_3$  becomes smaller at high annealing temperatures, the heat of

3/076/62/036/009/002/011 B1::1/B102

Study of the surface ...

wetting calculated per g of Al<sub>2</sub>O<sub>3</sub> reaches a maximum for Al<sub>2</sub>O<sub>3</sub> heated at 50000. The curve for heat of retting (Q, erg/cm<sup>3</sup>) versus structural vater (mole/m<sup>2</sup>) shows the following sections: (1) Increase of Q after thermal treatment of Al<sub>2</sub>O<sub>3</sub> at 30 - 150°C owing to removal of the adsorbed H<sub>2</sub>O<sub>1</sub> (2) treatment of Al<sub>2</sub>O<sub>3</sub> at 70 - 200°C in spite of dehydration of the bayerit in the bulk of Al<sub>2</sub>O<sub>3</sub>; (3) Q increases at 200 - 500°C owing to dehydration of the bulk of Al<sub>2</sub>O<sub>3</sub>; (4) sharp increase of Q between 500 and 700°C, although the Al<sub>2</sub>O<sub>3</sub> surfaces (4) sharp increase of Q between 500 and 700°C, although the content of structural water changes only little in this range owing to formation of Al<sub>2</sub>O<sub>3</sub>; (5) increase of Q at 800-900°C owing to formation of K, 8, 9, and 2-Al<sub>2</sub>O<sub>3</sub> (corrundum). The infrared spectrum of deuterated Al<sub>2</sub>O<sub>3</sub> showed a broad 2630 cm<sup>-1</sup> band which disappeared at 400°C (interacting Observations), a narrow band at 2755 cm<sup>-1</sup> (free, non-interacting Observations), and a narrow 2710 cm<sup>-1</sup> band (weakly bound Observations). For gibbsite, maximum hydration was calculated to be ~22µmole/m<sup>2</sup>; for the (0001) face of corundum, the hydration accounts to 12.7 µmole/m<sup>2</sup>. The coordination sphere of the Al Card 2/3

Study of the surface ...

3/076/62/036/009/002/011 B101/B102

surface atoms which is not fully occupied after the thermal dehydration is filled up by water or alcohols with formation of hydrate or alcoholates, mal treatment of Al<sub>2</sub>O<sub>3</sub> at high temperature. There are 4 figures and 2

ASJOCIATION:

Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova, Fizicneskiy i khimicheskiy fakul'tety (Moscow State University imeni M. V. Lomonosov, Physical and Chemical Departments)

SUBMITTED:

November 1, 1960

Card 3/3

TOPCHIYEVA, K.V.; ANTIPINA, T.V.; LI KHE-SUYAN'; LEONT'YEV, Ye.A.

Formation of the porous structure of aluminosilicate catalysts subjected to the action of surface-active agents. Kin.i kat. 2 no.6:887-893 N-D '61. (MIRA 14:12)

1. Moskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova, khimicheskiy fakul'tet.

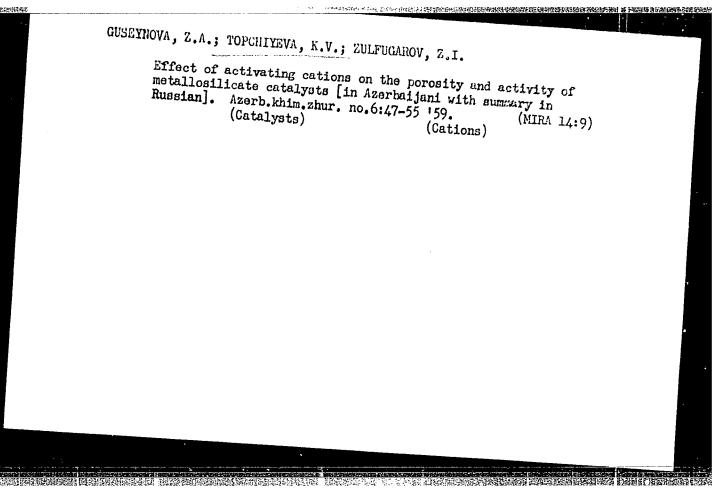
(Aluminosilicates) (Surface-active agents)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6"

TOPCHIYEVA, K.V.; PLANOVSKAYA, I.P.

Relationship between the extent of gaseous phase mixing in a fluidized bed and the flow rate and height of the catalyst layer. Dokl. AN SSSR 141 no.3:679-682 N '61. (MIRA 14:11)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova. Predstavleno akademikom S.I. Vol!fkovichem.
(Fluidization) (Catalysts)



TOPCHIYEVA, K.V.; VEN'YAMINOV, S.A.

Hydrochlorination of acetylene on aluminum oxide. Kin.i kat. 3 no.1:118-122 '62. (MIRA 15:3)

S/204/61/001/006/001/004 E075/E436

AUTHORS:

Topchiyeva, K.V., Sharayev, O.K., Perel'man, A.I.,

TITLE:

Some data on the polymerization of ethylene on chromia

PERIODICAL: Neftekhimiya, v.1, no.6, 1961, 780-785 The object of the work was to continue the investigation of ethylene polymerization process on chromia catalyst in order to elucidate the nature of the catalytic activity. The chromia catalyst was deposited on alumino-silicate obtained from silica gel covered with 3% wt of Al203. One portion of the catalyst was activated in N (dry air stream) for 4 hours at 500°C. portion was activated under vacuum at 350°C for 4 hours. The other 1.96% wt for the catalysts activated under vacuum and in N The quantities of  $Cr^{6+}$  were 1.25 and respectively. Experiments were carried out at several temperatures between 40 and 135°C. Ethylene was fed into reactor at the rate of 40 ml/min and each experiment lasted 40 min. Activity of the catalysts was obtained from the increases in their

Some data on the polymerization ... \$/204/61/001/006/001/004

weight due to deposition of polymer. evolved during the process was measured to observe the progress of the polymerization. It was assumed that the rise of the catalyst temperature T is proportional to the heat evolved and, consequently, to the reaction rate, obtained corresponded to the heat of adsorption. The small initial heating by an induction period and the main heating effect due to the polymerization. It was followed The heating curve rose exponentially, passed through a maximum and then fell as the reaction rate decreased. The length of the induction period increased (from about 2 to 20 min) with the decreasing temperature of reaction. induction period disappeared when the catalyst was activated with ethylene instead of nitrogen at 500°C. The authors concluded that the formation of active surface on chromia catalyst was due to its interaction with ethylene. was governed by the time of activation of the surface. The length of the induction periods authors postulated that ethylene reduced chromium in the catalyst The catalyst activated in air operated without the induction period because such a catalyst could be

### "APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001756310011-6 Comments of the second control of the second

Some data on the polymerization ... s/204/61/001/006/001/004 E075/E436 easily reduced,

This did not apply to the catalyst activated under vacuum (containing chromium chromates) which was much more difficult to reduce. The formation of the reduced form of chromium was confirmed by electron pararesonance spectra.

ASSOCIATION: Institut neftekhimicheskogo sinteza AN SSSR

Moskovskiy gosudarstvennyy universitet Kafedra fizicheskoy khimii

(Institute of Petrochemical Synthesis AS USSR Moscow State University

Department of Physical Chemistry)

SUBMITTED:

October 12, 1961

Card 3/3

SMIRNOVA, I.V.; KUBASOV, A.A.; TOPCHIYEVA, K.V.

Heat of wetting aluminum oxides by benzene, cyclchexane, and cyclohexene solutions in n-heptane. Dokl. AN SSSR 139 no.1:

(MIRA 14:7)

1. Moskovskiy gosudarstvennyy universitet im. M,V. Lomonosova.

(Aluminum oxide) (Heat of wetting)

TOPCHIYEVA, K.V.; ZEN'KOVICH, I.A.; BUKANAYEVA, F.M.

Catalytic activity of rare earth oxides deposited on silica in reactions involving the decomposition of alcohol. Vest. Mosk. un. Ser. 2:

(MIRA 1/:/)

1. Kafedra fizicheskoy khimii Moskovskogo universiteta.
(Rare earth oxides) (Dehydration (Chemistry))

### TOPCHIYEVA, K.V.

Nature of the acidity of aluminosilicate cracking catalysts. Probl. kin. 1 kat. 10:247-254 '60. (MIRA 14:5)

1. Khimicheskiy fakuL'tet Moskovskogo Gosudarstvennogo universiteta.
(Aluminosilicates) (Catalysts)

S/189/60/000/005/001/006 B110/217

AUTHORS:

Topchiyeva, K. V., Zen'kovich, I. A., Bukanayeva, F. M.

TITLE:

Effect exerted by the addition of rare earth oxides upon the catalytic properties of some oxidizing catalysts in hydro-

carbon reactions

PERIODICAL:

Vestnik Moskovskogo universiteta. Seriya 2, khimiya, no. 5,

1960, 3-5

Rare earths (Sm203; Nd203) are good dehydrogenating and cyclizing catalysts for paraffins and cycloparaffins, the activity of which is greatly increased by mixing with Al203. The authors aimed at obtaining a catalyst with bifunctional action (rare earth component for dehydrogenation) by adding rare earth oxides to aluminum silicate. The most active aluminum silicate  $(30\% \text{ Al}_20_3; 70\% \text{ SiO}_2)$  with admixtures of 5% of the total weight of  $\text{La}_20_3$ ;  $^{\mathrm{Nd}}_{2}^{0}_{3}$ ;  $^{\mathrm{Sm}}_{2}^{0}_{3}$ ;  $^{\mathrm{Pr}}_{2}^{0}_{3}$ ;  $^{\mathrm{Y}}_{2}^{0}_{3}$ ;  $^{\mathrm{Yb}}_{2}^{0}_{3}$ , was tested. Al(OH)<sub>3</sub>, silica gel, and rare earth hydroxide were mixed and activated in the  $N_2$  current at  $550\,^{\circ}\text{C}$  to pro-Card 1/5

Effect exerted by ...

S/189/60/000/005/001/006 B110/B217

duce the catalysts. Each experiment was followed by reactiviation in the air current at 500-550°C. Cumene cracking was studied at 450°C and a volume rate of 1 ml/ml·hr. When 5% oxide were added, the cracking ratio, mole of the separated gas: mole of passed through cumene decreased from 45% to 35%. In-octane was also investigated at 500°C and a volume rate of 0.65 ml/ml hr. The ratio, gas weight: weight of the passed through n-octane decreased by 20 mole%, with gas- and catalyzate composition remaining unchanged after analysis by means of BIM(VTI) apparatus. 5% Nd<sub>2</sub>0<sub>3</sub> admixture at 320°C, H<sub>2</sub> pressure = 24 atm., volume rate, 1 ml/ml·hr resulted at unchanged composition of the catalyzate in a decrease of cracking by ×7 mole%. This reduction of activity is due to a contamination of the acid aluminum silicate centers by the strongly basic hydroxides of the rare earths and partial destruction of the aluminum silicate structure. Also the catalysts: 95% Al<sub>2</sub>0<sub>3</sub>: 5% Pr<sub>2</sub>0<sub>3</sub>; 80% Al<sub>2</sub>0<sub>3</sub>: 5% Yb<sub>2</sub>0<sub>3</sub>; 95% Al<sub>2</sub>0<sub>3</sub>: 5% Sm<sub>2</sub>0<sub>3</sub>; 80% Al<sub>2</sub>0<sub>3</sub>: 20% La<sub>2</sub>0<sub>3</sub>; 80% Al<sub>2</sub>0<sub>3</sub>: 20% Pr<sub>2</sub>0<sub>3</sub>, with n-octane at 500-545°C and a volume rate of 0.64-0.16 ml/ml·hr, resulted in no increase of activity. The increase of cracking by ≈6-10% obtained with 80% Al<sub>2</sub>0<sub>3</sub>: 20% Pr<sub>2</sub>0<sub>3</sub> at a volume rate of Card 2/5

Effect exerted by ...

\$/189/60/000/005/001/006 B110/B217

0.16 ml/ml hr is due to the hydrogenation properties of  $Pr_2O_3$ . The results the authors obtained with the following catalysts: 85%  ${
m Al}_2{
m O}_3$ : 15%  ${
m Me}_2{
m O}_3$ (Me = Nd, Sm) were in complete disagreement with those of V. I. Komarewsky (Ref. 1: Industr. and Engng. chem., 49, No. 2, 264-265, 1957). The experiment made by this researcher with heptane and 85% Al203 with 15% Nd203 was repeated, the catalyst being produced by his method of mixing and coprecipitation. The calculated amount of highly acid 0.39 M Nd(NO3) was added to 0.725 M sedium aluminate solution. The catalyst was activated at 550°C in the  $\mathrm{N}_2$  current. No increase of activity as compared to pure  $\mathrm{Al}_2\mathrm{O}_3$  was established. Possibly, Komerewsky prepared his mixing catalysts in a different way, or he compared their activity with that of the rare earth exide and thought that Al203 was inactive. The higher activity of his catalysts may also be due to Al203 which, according to its way of preparation, may also have dehydrogenating properties (Table). There are 1 table and 3 references: 1 Soviet-bloc and 2 non-Soviet-bloc. The reference to Englishlanguage publications reads as follows: Ref. 2: Ciapetta F. G., Hunter J. Card 3/5

Effect exerted by ...

**3/189/60/000/005/001/006 B110/B217** 

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B. Industr. and Engng. chem., 45, 147-55, 1953.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lamanosava, Kafedra fizisheskoy khimii (Moscow State University imeni M. V. Lamanosav Department of Physical Chemistry)

SUBMITTED: July 14, 1959

Legend to the Table: The conversion of n-heptane at 525°C on the mixing catalyst, 85% Al<sub>2</sub>O<sub>3</sub>: 15% Nd<sub>2</sub>O<sub>3</sub>; 1) catalyst: volume rate ml/ml·hr; 2) thermal cracking 4.85 ml/hr; 3) coprecipitation method; 4) mixing method; 5) data by Komarewsky; 6) bulk factor of the catalyst, ml; 7) yield, wt%; 8) of gas; 9) of catalyst; 10) losses; 11) gas composition, vol%; 12) paraffins; 13) and 14) olefins; 15) aromatic components; 16) catalyzate composition, wt%.

Card 4/5

Effect	exerted by						s/18 B110	39/6 0/B2	0/00 17	0/00	5/00	1/00	6	
	Превращение и-гептана при $525^{\circ}$ на смещанном, катализаторе состава $85^{\circ}{}_{i0}$ Al <sub>2</sub> O <sub>3</sub> : $15^{\circ}{}_{i0}$ Nd <sub>2</sub> O <sub>3</sub> : $15^{\circ}{}_{i0}$													
	, 1		Выход. вес. %		Состав газа, 1106 гени. %			Состав ката- лизата, всс. 16 %		· ' .				
	<ul> <li>1 Катализатор: объемная скорость, мл/мл⋅час</li> </ul>	Насыпной о	r338 S	у катализата	потери 10	Нэ	парафийы	олефину.	4 k oreфints	45. арочатнка				
	2 Терынческий крекинг 4,58 мл/час .	-	15,6	85,0	O	1,00	85,8	13,2	1-2					
	Al <sub>2</sub> O <sub>3</sub> 0,15	30	22,6	72,3	5,1	18,1	71,5	10,4	3-4	2				
	3 (Метод соосаждения) 0,15 85% Al <sub>2</sub> O <sub>3</sub> :15% Nd <sub>2</sub> O <sub>3</sub>	30	17,2	70,2	12,6	15,8	73,2	11,0	5-7	2	,		•	/
	4 (Метод смещения) 0.15 85 <sup>9</sup> / <sub>0</sub> Al <sub>2</sub> O <sub>3</sub> :15 <sup>9</sup> / <sub>0</sub> Nd <sub>2</sub> O <sub>3</sub>	30	21,9	62,8	12,3	18,5	68,9	12,6	3					1
	5 (Данные Комаренского) 0.15 85% Al <sub>2</sub> O <sub>3</sub> :15% Nd <sub>2</sub> O <sub>3</sub>	30		71,8	t .	64,8	22,	12,3	10,3	21				
Card 5/	'5													

TOPCHIYEVA, K.V.; HAHBAYEVA, A.M.; SPOZHAKIKA, A.A.

Effect of hydrogen chloride on the catalytic properties of aluminum oxide in the reaction of cracking. Vest. Mosk. un. Ser. 2: Whir. 15 no.6:10-14 M-D '60. (MIRA 14:2)

1. Kafedra fizicheskoy khimii Moskovskogo universiteta.
(Hydrochloric acid) (Alumina)
(Cracking process)

Studying the kinetics of the cracking of cumene on a fluidized

aluminosilicate catalyst. Vest. Mosk.un. Ser. mat., mekh., astron. fiz., khim. 14 no.3:151-157 '59. (MIRA 13:5)

1. Kafedra fizicheskoy khimii Moskovskogo gosudarstvennogo universiteta.

(Cumene) (Cracking process)

TOPCHIYEVA, K.V.; ZEN!KOVICH, I.A.; BUKANAYEVA, F.H.

Effect of rare earth oxide impurities on the catalytic properties of some oxide catalysts in reactions of hydrocartons. Vest. Mosk. un. Ser. 2: Khim. 15 no.5:3-5 S-0 '60. (MIRA 13:11)

1. Moskovskiy gosudarstvennyy universitet, kafedra fizicheskov khimii.

(Rare earth oxides) (Catalysts)

S/189/60/000/002/002/008/XX B017/B067

AUTHORS:

Topchiyeva, K. V., Antipina, T. V., and Khe-Suyan', Li

TITLE:

Effect of the Pore Radius and Other Structural Characteristics of Oxidic Catalysts on the Parameters of

the Course of Heterogeneous Catalytic Processes, Communication I. Production of Aluminum Silicate Catalysts With Different Structural Characteristics

PERIODICAL:

Vestnik Moskovskogo universiteta. Seriya 2, khimiya, 1960,

No. 2, pp. 13 - 21

TEXT: Aluminum silicate catalysts of equal chemical composition but different structural characteristics were produced in experimental series. The adsorption- and structural properties of the catalysts were studied in dependence on the degree of displacement of intermicellar water by isobutyl alcohol and cumene. Intermicellar water was displaced by isobutyl alcohol and cumene in the apparatus shown in Fig.1. Aluminum silicate catalysts of the composition 12% Al<sub>2</sub>O<sub>3</sub> and 88% SiO<sub>2</sub> were

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Effect of the Pore Radius and Other S/189/60/000/002/002/002/008/IX Structural Characteristics of Oxidic B017/B067 Catalysts on the Parameters of the Course of Heterogeneous Catalytic Processes. Communication I. Production of Aluminum Silicate Catalysts With Different Structural Characteristics

produced by the method of GrozNII (Groznyy Scientific Research Institute). The structural characteristics of aluminum silicate catalysts are given in a Table. Two series of catalysts were produced: 1) Series with the intermicellar water being displaced by butyl alcohol at 35, 51, 81, 95, 97, and 100%; 2) series with the intermicellar water being displaced by cumene at 34, 63, 90, and 96%. The adsorptive properties and structural characteristics of the catalysts produced were calculated from the isotherms of adsorption of methyl alcohol vapor at 20°C. Figs. 2 and 4 graphically show the adsorption isotherms of methyl alcohol vapor, the distribution of pore volumes and pore radii, and the dependence of the structural characteristics on the degree of displacement of water in the catalysts of the first series. Figs. 5-7 show the same curves for the catalysts of the second series, in which water was displaced by cumene. It was observed that the chemical nature and properties of the organic solvents, which displace the water from the catalysts, influence the structure of the catalysts. An aluminum silicate catalyst with

Card 2/3

Effect of the Pore Radius and Other Structural Characteristics of Oxidic s/189/60/000/002/002/008/xx Catalysts on the Parameters of the Course of Heterogeneous Catalytic Processes. Communication I. Production of Aluminum Silicate Catalysts With Different Structural Characteristics

a specific surface ( $S_{\rm BET} = 450~{\rm m}^2/{\rm g}$ ) was produced. The authors mention Professor I. Ye. Neymark and A. V. Kiselev. There are 7 figures, 1 table, and 6 Soviet references.

ASSOCIATION: Kafedra fizicheskoy khimii (Chair of Physical Chemistry)

SUBMITTED: October 2, 1959

Card 3/3

S/189/60/000/002/003/008/XX B017/B067

AUTHORS:

Topchiyeva, K. V. and Moskovskaya, I. F.

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TITLE:

Chemosorption of Hydrogen on Aluminum Silicates and

Aluminum and Silicon Oxides \_ v

PERIODICAL:

Vestnik Moskovskogo universiteta. Seriya 2, khimiya, 1960,

No. 2, pp. 22 - 27

TEXT: The authors studied the adsorption of hydrogen on aluminum silicate catalysts of different compositions, and on aluminum and silicon oxides at different temperatures and pressures. The apparatus for determining the hydrogen adsorption in vacuo is schematically shown in Fig.1. The following catalysts were used for adsorption experiments: 1) Aluminum silicates with a percent composition of 30/70, 50/50, 80/20 Al $_20_3/\sin 2$ , and 2) pure aluminum and silicon oxides which were

produced by a method described in Ref.13. The adsorptions were made in a temperature range of from 23 to 600°C. The course of adsorption on aluminum silicate catalysts at different temperatures with a percent

Card 1/2

Chemosorption of Hydrogen on Aluminum Silicates and Aluminum and Silicon Oxides

S/189/60/000/002/003/008/XX B017/B067

composition of 30/70 and 80/20 Al<sub>2</sub>O<sub>3</sub>/SiO<sub>2</sub> is graphically shown in Fig.2. An increased adsorption of hydrogen on aluminum silicate catalysts occurs at 550-600°C. For aluminum oxide, this region of increased hydrogen adsorption is at 300-600°C, for silicon oxide at 400-600°C. Adsorption energy and adsorption heat were determined. Chemosorption of hydrogen on catalysts occurs in newly formed, unstable, active centers which are destroyed on a regeneration of the catalysts. The amount of adsorbed hydrogen is low and covers less than 1% of the specific surface of the catalysts. There are 5 figures, 1 table, and 14 references: 7 Soviet, 3 US, and 3 German.

ASSOCIATION: Kafedra fizicheskoy khimii (Chair of Physical Chemistry)

SUBMITTED: March 16, 1959

Card 2/2

TOPCHIYEVA, K.V.; TAKHTAROVA, G.N.; FOMINA, A.I.

Vapor-phase esterification of aromatic acids with atters on exide catalysts. Neftekhimita 2 no.5:744-749 S-0 '62.' (MIRA 16:1)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimicheskiy fakul'tet.
(Acids, Organic) (Esterification) (Ethers)

S/204/63/003/001/006/013 E075/E436

AUTHORS:

Smirnova, I.V., Karpukhina, G.V., Topchiyeva, K.V.

TITLE:

Adsorption of allylbenzene and allylcyclohexane on

chromia catalyst

PERIODICAL: Neftekhimiya, v.3, no.1, 1963, 71-73

The adsorption from n-heptane of the two hydrocarbons on Croz was studied to gain an insight into the mechanism of the polymerization of unsaturated hydrocarbons. prepared by a previously described method (A.V. Topchiyev et al. Dokl. AN SSSR, v.130, 1960, 344) and had the surface area of There were no catalytic reactions taking place during the experiments. The adsorption isotherms were determined at 20°C by interferometry. Allylbenzene was shown to occupy an area on the catalyst similar to that occupied by benzene on silica gel or alumina. Allyl groups were apparently above the level of the adsorbed benzene nuclei making the adsorbed mono-layer relatively thick and not in contact with the catalyst surface. The molecules of adsorbed allylcyclohexane occupied much larger area, the allyl groups being in direct contact with the surface.

5/204/63/003/001/006/013 E075/E436

Adsorption of allylbenzene ...

Because of this fact, it is considered that allylcyclohexane and other allylnaphthenes should polymerize more easily than allylbenzene. There is 1 figure.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im.

M.V.Lomonosova (Moscow State University imeni

M.V.Lomonosov)

July 7, 1962 SUBMITTED:

TPCHIYEVA, K.V., ANTIPINA, T.V., LI KHE-SUYAN¹

Effect of the size of pore radii and other structural characteristics of oxide catalysts upon the parameters of the course of heterogeneous catalytic processes. Vest. Mosk. un. Ser. 2: khim. 15 no.2:13-21 Mr-Ap '60.

1. Kafedra fizicheskoy khimii Moskovskogo universiteta.

(Catalysts) (Aluminosilicates)

TOPCHIYEVA, K.V.; DANILOVA, N.A.; MAKAROVA, A.M.

Investigating the effect of high temperature and water vapor on the structure and activity of magnesium silicate catalysts.

Azerb,khim.shur. no.2:85-91 159. (MIRA 13:6)

(Magnesium silicate) (Catalysis)

SMIRNOVA, V.Ye.; TOPCHIYEVA, K.V.; ZUL'FUGAROV, Z.G.

Effect of the chemical composition, the pH of the synthesis medium, and the nature of the initial sols on the activity of medium, and the nature of the initial sols on the activity of aluminosilicate catalysts. Azerb.khim.zhur. no.1:83-95 (MIRA 13:6)

'59. (Aluminosilicates) (Catalysis)

FROST, Andrey Vladimirovich, prof. [deceased]; Prinimali uchastiye:

BUSHMAKIN, I.N.; VYEDENSKIY, A.A.; GRYAZNOV, V.M.; DEMZHT'YEVA,

M.I.: DINTSES, A.I.; DOBROHRAVOV, R.K.; ZHARKOVA, V.R.; ZHERKO,

A.V.; IPAT'YEV, V.N.; KVYATKOVSKIY, D.A.; KOROBOV, V.V.; MOOR,

V.G.; NEMTSOV, M.S.; RAKOVSKIY, A.V.; REMIZ, Ye.K.; RUDKOVSKIY,

D.M.; RYSAKOV, M.V.; SEREBRYAKOVA, Ye.K.; STEPUKHOVICH, A.D.;

STRIGALEVA, N.V.; TATEVSKIY, V.M.; TILICHEYEV, M.D.; TRIFEL',

A.G.: FROST, O.I.; SHILYAYEVA, L.V.; SHCHEKIN, V.V., DOLGOPOLOV,

N.N., SOSTAVITEL'; GERASIMOV, Ya.I., otv.red.; SMIRNOVA, I.V.; red.;

TOPCHIYEVA, K.V.; YASTREBOV, V.V., red.; KONDRASHKOVA, S.F., red.

izd-va; LAZAREVA, L.V., tekhn.red.

[Selected scientific works] Izbrannye nauchnye trudy. Moskva, Izd-vo Mosk.univ., 1960. 512 p. (MIRA 13:5)

1. Chlen-korrespondent AN SSSR (for Gerasimov). (Chemistry, Physical and theoretical)

TOPCHIYEVA, K.V., MOSKOVSKAYA, I.F.

Chemisorption of hydrogen on aluminosilicates and aluminum and

silicon oxides. Vest. Mosk. un. Ser. 2: khim. 15 no.2:22-27 Mr-Ap , 60. (MIRA 13:6)

1. Kafedra fizicheskoy khimii Moskovskogo universiteta.
(Hydrogen) (Aluminosilicates) (Alumina) (Silica)

TOPCHIYEVA, K.V.; ROMANOVSKIY, B.V.; TIMOSHENKO, V.I.

Kinetics of heterogeneous catalytic reactions studied by the circulation method. Part 2: Cumene cracking over aluminosilicate catalysts. Kin.i kat. 6 no.3:4/71-475 My-Je '65.

(MIRA 18:10)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova, khimicheskiy fakulitet,

TOPCHIYEVA, K.V.; STETSENKO, V.Ya.

New catalytic properties of yttrium oxide. Kin. i kst. 6 no.4:751

JI-Ag 165. (MIRA 18:9)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova, Khimicheskiy fakul $^{\dagger}$ tet.

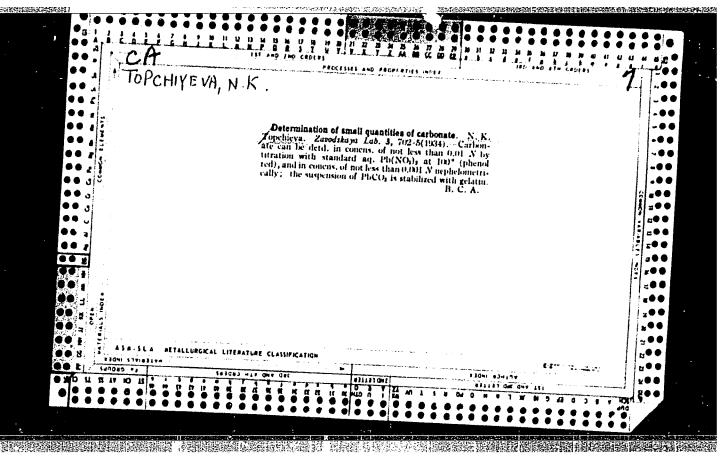
KIRICHENKO, A.G., inzh.; TURCHIYEVA, M.V., inch.; NEVYOROV, M.\*., inzh.;
PANTELYAT, G.S., inzh.

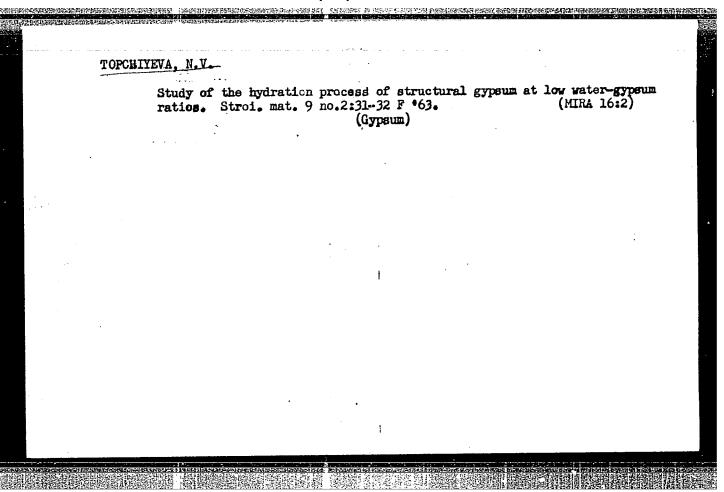
Biochemical consumption of oxygen by the waste waters of Kharkov. Vod. i san. tekh. no.4:12-14 Ap \*65.

(MTRA 19:1)

MEMINA, A.T.; TOPCHIYEVA, M.V.

Operations of primary horizontal clarifiers at the Kharkov biological station. Vod. i san. tekh. no.9:3-7 S '58. (MIRA 11:10) (Kharkev--Water--Purification)





TOPCHIYEVA, N.V.

Regulating the processes of setting and hardening of plasters by the two-stage solution method[with summary in English]. Dop.AN URSR no.3:363-367 '61. (MIRA 14:3)

l. Nauchno-issledovatel'skiy institut stroitel'nykh materialov i sooruzheniy Akademii stroitel'stva i arkhitektury USSR. Predstavleno akademikom AN USSR P.P.Budnikovym.

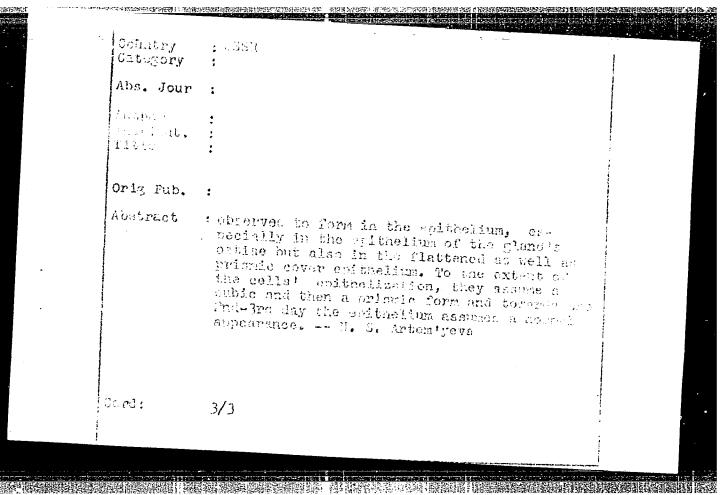
(Plaster)

Two-stage method for making gypsum mortars. Stroi.mat. 6 no.5: 35-36 My '60. (MHA 13:7)

(Gypsum)

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Country	: USSR
abs. Jour	:
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Cris Pub.	igurophic manifestations in underlaying tismes. The epithelization of the macesa is realized by moving in of the epithelial layer from the borders of the defect, and here, a flattening of the prismatic cells which are situated chiefly in ostiae of the glands, is observed. During the first 24 hours after the observed. The place and amitoses are not observed trauma, mitoses and amitoses are not observed to take place in epithelial cells. After 24 hours a significant number of mitoses are
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TOPCHIYEVA, O.I. (Leningrad, 49, ul. L.Chaykinoy, d.21, kv.24)

Regeneration of the uterine epithelium in superficial mechanical injury of the mucous membrane. Arkh.anat.gist. i embr. 33 no.4: 49-54 O-D '56. (MLRA 10:4)

1. Is otdels patologicheskoyanatomii (zaveduyushchiy - akademik N.N.Anichkov) Instituta eksperimental noy meditsiny AMN SSSR i kafedry akusheratva i ginekologii (zaveduyushchiy - professor I.I. Iakovlev) Leningradskogo meditsinskogo instituta im. akademika I.P. Pavlova.

(BEDOMETRIUM, wounds and inj. regeneration process, histol.)

(REGENERATION endometrium, after superficial inj., histol. aspects)